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         MAR 11
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         MAR 11
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         MAR 23
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         MAR 30
                 IMSPATENTS reloaded and enhanced
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         APR 03
                 CAS coverage of exemplified prophetic substances
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NEWS 23
         APR 24 CA/CAplus now has more comprehensive patent assignee
                 information
NEWS 24
         APR 26 USPATFULL and USPAT2 enhanced with patent
                 assignment/reassignment information
NEWS 25 APR 28
                 CAS patent authority coverage expanded
                 ENCOMPLIT/ENCOMPLIT2 search fields enhanced
NEWS 26
         APR 28
NEWS 27 APR 28 Limits doubled for structure searching in CAS
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NEWS 29 MAY 11 STN on the Web enhanced

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NEWS EXPRESS MAY 08 09 CURRENT WINDOWS VERSION IS V8.4, AND CURRENT DISCOVER FILE IS DATED 06 APRIL 2009.

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chain nodes :
1 2 17 26 28
ring nodes :
3 \quad 4 \quad 5 \quad 6 \quad 7 \quad 8 \quad 9 \quad 10 \quad 11 \quad 12 \quad 13 \quad 19 \quad 20 \quad 21 \quad 22 \quad 23 \quad 24
chain bonds :
1-2 1-26 11-17 17-19 26-28
ring bonds :
3-4 3-7 4-5 5-6 6-7 8-9 8-13 9-10 10-11 11-12 12-13 19-20 19-24 20-21
21-22 22-23 23-24
exact/norm bonds :
1-2 1-26 3-4 3-7 11-17 17-19 26-28
exact bonds :
4-5 5-6 6-7
normalized bonds :
8-9 8-13 9-10 10-11 11-12 12-13 19-20 19-24 20-21 21-22 22-23 23-24
isolated ring systems :
containing 3 : 19 :
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G1:0,S

G2:H,Ak

Match level:

1:CLASS 2:CLASS 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 17:CLASS 19:Atom 20:Atom 21:Atom 22:Atom 23:Atom 24:Atom 26:CLASS 28:CLASS

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2 ANSWERS

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L22 SEA SSS FUL L1

L3 1 L2

=> d ibib abs hitstr 1-YOU HAVE REQUESTED DATA FROM 1 ANSWERS - CONTINUE? Y/(N):y

ANSWER 1 OF 1 CAPLUS COPYRIGHT 2009 ACS on STN T.3 ACCESSION NUMBER: 2005:469894 CAPLUS Full-text DOCUMENT NUMBER: 143:7592

TITLE:

Preparation of arylpyrrolecarboxamides as Raf kinase

inhibitors for treatment of tumors.

INVENTOR(S): Finsinger, Dirk; Buchstaller, Hans-Peter; Burgdorf,

Lars; Wiesner, Matthias; Amendt, Christiane; Grell,

Matthias; Sirrenberg, Christian; Zenke, Frank

PATENT ASSIGNEE(S): Merck Patent G.m.b.H., Germany

SOURCE: Ger. Offen., 32 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA'	PATENT NO.							DATE		APPLICATION NO.					DATE			
DE	E 10354060						20050602		DE 2003-10354060					20031119				
							20050602		AU 2004-291255					20041026				
CA	CA 2546334				A1 2005060			0602	CA 2004-2546334					20041026				
WO	2005049603				A1 2005			0602	WO 2004-EP12076						20041026			
	W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB	BG,	BR,	BW,	BY,	BZ,	CA,	CH,	
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ	EC,	EE,	EG,	ES,	FΙ,	GB,	GD,	
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS	JP,	KE,	KG,	KP,	KR,	KΖ,	LC,	
		LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG	, MK,	MN,	MW,	MX,	MZ,	NA,	NΙ,	
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		ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US	, UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW	
	RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD	, SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	
		ΑZ,	BY,	KG,	KΖ,	MD,	RU,	ТJ,	TM,	ΑT	, BE,	BG,	CH,	CY,	CZ,	DE,	DK,	
		EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,	ΙT	LU,	MC,	NL,	PL,	PT,	RO,	SE,	
		SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM	I, GA,	GN,	GQ,	GW,	${ m ML}$,	MR,	ΝE,	
		SN,	TD,	ΤG														
EP	EP 1685125				A1 20060802			EP 2004-790859						20041026				
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR	R, IT,	LI,	LU,	NL,	SE,	MC,	PT,	
											CZ,		•					
CN 1882571																		
BR 2004016690				Α		20070130			BR 2004-16690					20041026				
JP 2007511553				Τ	T 20070510				JP 2006-540216				20041026					
IN 2006KN00936					A		2007	20070420 :			2006-		20060417					
MX 2006005478										MX 2006-5478								
KR 2006118492																		
US 20070149594				A1		20070628			US 2006-579825									
PRIORITY APPLN. INFO.:				.:							2003-							
										WO	2004-	EP12	076		W 2	0041	026	
OTHER SOURCE(S):					MAR:	PAT	143:	7592										

Z-N N XAr

GΙ

Title compds. [I; Ar = (substituted) Ph, naphthyl, biphenyl, heterocyclyl; X = 0, S, (CH2)n, CO, (CH2)nO, (CH2)nNH, etc.; n = 1-3; Y = 0, S, CHNO2, C(CN)2, NR4; R4 = H, cyano, OH, etc.; Z = Ar, ArXAr, CH2Ar, CH2ArXAr; Ar = (substituted) Ph], were prepared as Raf kinase inhibitors (no data). Thus, 4-

(PhCH2O)C6H4CH2CO2H, DMF, and POC13 were heated together at 70° for 4 h followed by cooling and addition of ice water and aqueous NaC1O4 to give 98% [2-(4-benzyloxyphenyl)-3-dimethylaminoallylidene]dimethylammonium perchlorate. This was refluxed 24 h with glycine Et ester hydrochloride in EtOH containing 20% NaOEt to give 91% Et 4-(4-benzyloxyphenyl)-1H-pyrrole-2-carboxylate. Hydrogenolysis of the latter in EtOAc over Pd/C gave 91% Et 4-(4-hydroxyphenyl)-1H-pyrrole-2-carboxylate. This was heated with 4-chloropyridine-2-carboxylic acid N-methylamide at 160° for 48 h to give 40% Et 4-[4-(2-methylcarbamoylpyridin-4-yloxy)phenyl]-1H-pyrrole-2-carboxylate. Saponification with 2N NaOH in EtOH at 60° for 16 h followed by acidification with HCl gave 85% free acid, which was stirred 48 h in DMF with 5-amino-2-chlorobenzotrifluoride, N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride, and 1-hydroxybenzotriazole hydrate to give 17% 4-[4-[5-(4-chloro-3-trifluoromethylphenylcarbamoyl)-1H-pyrrol-3-yl]phenoxy]pyridine-2-carboxylic acid N-methylamide.

IT 852455-40-4P 852455-41-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of arylpyrrolecarboxamides as Raf kinase inhibitors for treatment of tumors)

RN 852455-40-4 CAPLUS

CN 1H-Pyrrole-2-carboxylic acid, 4-[4-[[2-[(methylamino)carbonyl]-4-pyridinyl]oxy]phenyl]-, ethyl ester (CA INDEX NAME)

RN 852455-41-5 CAPLUS

CN 1H-Pyrrole-2-carboxylic acid, 4-[4-[[2-[(methylamino)carbonyl]-4-pyridinyl]oxy]phenyl]- (CA INDEX NAME)

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